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Complete Listing of ALL CLAIMS, WITH MARKINGS AND STATUS IDENTIFIERS (Currently amended claims showing deletions by strikethrough and additions by underlining)

1-2 (canceled)

3 (currently amended): A method of claim 2 decreasing body weight in a patient, wherein said method comprising administering a therapeutically effective amount of a somatostatin agonist is a somatostatin type-2 receptor agonist to said patient.

4 (canceled)

5 (currently amended): A <u>The</u> method of claim 3, wherein said somatostatin type-2 receptor agonist has a Ki of less than 2 nM for the somatostatin type-2 receptor.

6 (canceled)

7 (currently amended): A $\underline{\text{The}}$ method of claim $\frac{2}{3}$, wherein said somatostatin agonist is a somatostatin type-2 receptor selective agonist.

8 (canceled)

9 (currently amended): A <u>The</u> method of claim 7, wherein said somatostatin type-2 receptor selective agonist has a Ki for the somatostatin type-2 receptor that is at least 10 times less than the Ki for the somatostatin type-1, type-3, type-4, and type-5 receptors.

10-13 (canceled)

14 (currently amended): A <u>The</u> method of claim 3, wherein said patient is a non-insulin-dependent diabetic human.

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15 (canceled)

16 (currently amended): A <u>The</u> method of claim 5, wherein said patient is a non-insulin-dependent diabetic human.

17 (canceled)

18 (currently amended): A <u>The</u> method of claim 7, wherein said patient is a non-insulin-dependent diabetic human.

19 (canceled)

20 (currently amended): A <u>The</u> method of claim 9, wherein said patient is a non-insulin-dependent diabetic human.

21-22 (canceled)

23 (currently amended): A The method according claim 1 3 wherein the somatostatin agonist is H-D-ß-Nal-Cys-Tyr-D-Trp-Lys-Thr-Cys-Thr-NH2, H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-S-Nal-NH2, H-D-Phe-Cys-Tyr-D-Trp-Lys-Thr-Cys-S-Nal-NH2, H-D-ß-Nal-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH, H-D-Phe-Cys-Tyr-D-Trp-Lys-Thr-Pen-Thr-NH2, H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Pen-Thr-NH2, H-D-Phe-Cys-Tyr-D-Trp-Lys-Thr-Pen-Thr-OH, H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Pen-Thr-OH, H-Gly-Pen-Phe-D-Trp-Lys-Thr-Cys-Thr-OH, H-Phe-Pen-Tyr-D-Trp-Lys-Thr-Cys-Thr-OH, H-Phe-Pen-Phe-D-Trp-Lys-Thr-Pen-Thr-OH, H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-ol H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2, H-D-Trp-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH,, H-D-Trp-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,

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H-D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2,
H-D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Trp-NH,,
H-D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2,
Ac-D-Phe-Lys*-Tyr-D-Trp-Lys-Val-Asp-Thr-NH2 (an amide bridge formed
between Lys and Asp),
Ac-hArq(Et),-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-D-hArq(Et),-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-D-hArg(Bu)-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-D-hArg(Et),-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-L-hArq(Et),-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-D-hArg(CH2CF3)2-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-D-hArg(CH<sub>2</sub>CF<sub>3</sub>)<sub>2</sub>-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH<sub>2</sub>,
Ac-D-hArg(CH2CF3)2-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Phe-NH2,
Ac-D-hArg(CH2CF3)2-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NHEt,
Ac-L-hArg(CH2-CF3)2-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-D-hArg(CH<sub>2</sub>CF<sub>3</sub>)<sub>2</sub>-Gly-Cys-Phe-D-Trp-Lys(Me)-Thr-Cys-Thr-NH<sub>2</sub>,
Ac-D-hArg(CH<sub>2</sub>CF<sub>3</sub>)<sub>2</sub>-Gly-Cys-Phe-D-Trp-Lys(Me)-Thr-Cys-Thr-NHEt,
Ac-hArq(CH<sub>2</sub>, hexyl)-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH<sub>2</sub>,
H-hArg(hexyl<sub>2</sub>)-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH<sub>2</sub>,
Ac-D-hArg(Et)2-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NHEt,
Ac-D-hArg(Et)2-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Phe-NH2,
Propionyl-D-hArg(Et)<sub>2</sub>-Gly-Cys-Phe-D-Trp-Lys(iPr)-Thr-Cys-Thr-NH<sub>2</sub>,
Ac-D-\( S-Nal-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Gly-hArg (Et)_2-NH_2,
Ac-D-Lys(iPr)-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-D-hArg(CH,CF,),-D-hArg(CH,CF,),-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-
Thr-NH2,
Ac-D-hArg(CH<sub>2</sub>CF<sub>3</sub>)<sub>2</sub>-D-hArg(CH<sub>2</sub>CF<sub>3</sub>)<sub>2</sub>-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-
Ac-D-hArq(Et),-D-hArq(Et),-Gly-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NH2,
Ac-Cys-Lys-Asn-4-Cl-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-Ser-D-Cys-NH2,
H-Bmp-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2,
H-Bmp-Tyr-D-Trp-Lys-Val-Cys-Phe-NH2,
H-Bmp-Tyr-D-Trp-Lys-Val-Cys-p-Cl-Phe-NH<sub>2</sub>,
H-Bmp-Tyr-D-Trp-Lys-Val-Cys-ß-Nal-NH2,
H-D-S-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2,
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H-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH2,
H-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-$-Nal-NH2,
H-pentafluoro-D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2,
Ac-D-ß-Nal-Cys-pentafluoro-Phe-D-Trp-Lys-Val-Cys-Thr-NH,,
H-D-S-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-S-Nal-NH2,
H-D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-\(\mathbb{G}\)-Nal-NH,
H-D-ß-Nal-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH2,
H-D-p-Cl-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH2,
Ac-D-p-Cl-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH,
H-D-Phe-Cys-ß-Nal-D-Trp-Lys-Val-Cys-Thr-NH2,
H-D-Phe-Cys-Tyr-D-Trp-Lys-Cys-Thr-NH2,
cyclo(Pro-Phe-D-Trp-N-Me-Lys-Thr-Phe),
cyclo(Pro-Phe-D-Trp-N-Me-Lys-Thr-Phe),
cyclo(Pro-Phe-D-Trp-Lys-Thr-N-Me-Phe),
cyclo(N-Me-Ala-Tyr-D-Trp-Lys-Thr-Phe),
cyclo(Pro-Tyr-D-Trp-Lys-Thr-Phe),
cyclo(Pro-Phe-D-Trp-Lys-Thr-Phe),
cyclo(Pro-Phe-L-Trp-Lys-Thr-Phe)(SEQ ID NO:1),
cyclo(Pro-Phe-D-Trp(F)-Lys-Thr-Phe),
cyclo(Pro-Phe-Trp(F)-Lys-Thr-Phe) (SEQ ID NO:2),
cyclo(Pro-Phe-D-Trp-Lys-Ser-Phe),
cyclo(Pro-Phe-D-Trp-Lys-Thr-p-Cl-Phe),
cyclo(D-Ala-N-Me-D-Phe-D-Thr-D-Lys-Trp-D-Phe),
cyclo(D-Ala-N-Me-D-Phe-D-Val-Lys-D-Trp-D-Phe),
cyclo(D-Ala-N-Me-D-Phe-D-Thr-Lys-D-Trp-D-Phe),
cyclo(D-Abu-N-Me-D-Phe-D-Val-Lys-D-Trp-D-Tyr),
cyclo(Pro-Tyr-D-Trp-t-4-AchxAla-Thr-Phe),
cyclo(Pro-Phe-D-Trp-t-4-AchxAla-Thr-Phe),
cyclo(N-Me-Ala-Tyr-D-Trp-Lys-Val-Phe),
cyclo(N-Me-Ala-Tyr-D-Trp-t-4-AchxAla-Thr-Phe),
cyclo (Pro-Tyr-D-Trp-4-Amphe-Thr-Phe),
cyclo (Pro-Phe-D-Trp-4-Amphe-Thr-Phe),
cyclo(N-Me-Ala-Tyr-D-Trp-4-Amphe-Thr-Phe),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-Gaba),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-Gaba-Gaba),
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cyclo(Asn-Phe-D-Trp-Lys-Thr-Phe),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-NH(CH2),CO),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-S-Ala),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-D-Glu)-OH,
cyclo(Phe-Phe-D-Trp-Lys-Thr-Phe),
cyclo(Phe-Phe-D-Trp-Lys-Thr-Phe-Gly),
cyclo(Phe-Phe-D-Trp-Lys-Thr-Phe-Gaba),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-Gly),
cyclo(Asn-Phe-Phe-D-Trp(F)-Lys-Thr-Phe-Gaba),
cyclo (Asn-Phe-Phe-D-Trp (NO2) -Lys-Thr-Phe-Gaba),
cyclo(Asn-Phe-Phe-Trp(Br)-Lys-Thr-Phe-Gaba) (SEQ ID NO:3),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Phe(I)-Gaba),
cyclo(Asn-Phe-Phe-D-Trp-Lys-Thr-Tyr(But)-Gaba),
cyclo(Bmp-Lys-Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-Pro-Cys)-OH,
cyclo(Bmp-Lys-Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-Pro-Cys)-OH,
cyclo(Bmp-Lys-Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-Tpo-Cys)-OH,
cyclo(Bmp-Lys-Asn-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-MeLeu-Cys)-OH,
cyclo(Phe-Phe-D-Trp-Lys-Thr-Phe-Phe-Gaba),
cyclo(Phe-Phe-D-Trp-Lys-Thr-Phe-D-Phe-Gaba),
cyclo(Phe-Phe-D-Trp(5F)-Lys-Thr-Phe-Phe-Gaba),
cyclo(Asn-Phe-Phe-D-Trp-Lys(Ac)-Thr-Phe-NH-(CH<sub>2</sub>)<sub>3</sub>-CO),
cyclo(Lys-Phe-Phe-D-Trp-Lys-Thr-Phe-Gaba),
cyclo(Lys-Phe-Phe-D-Trp-Lys-Thr-Phe-Gaba),
cyclo (Orn-Phe-Phe-D-Trp-Lys-Thr-Phe-Gaba),
H-Cys-Phe-Phe-D-Trp-Lys-Thr-Phe-Cys-NH, ,
H-Cys-Phe-Phe-D-Trp-Lys-Ser-Phe-Cys-NH2,
H-Cys-Phe-Tyr-D-Trp-Lys-Thr-Phe-Cys-NH2 or
H-Cys-Phe-Tyr(I)-D-Trp-Lys-Thr-Phe-Cys-NH<sub>2</sub>.
          24 (currently amended): A The method according to
claim 1 3 wherein the somatostatin agonist is
       ^{'}A^{1}-A^{2}-A^{3}-D-Trp-Lys-A^{6}-A^{7}-A^{8}-R_{3}
wherein
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A¹ is a D- or L- isomer of Ala, Leu, Ile, Val, Nle, Thr, Ser, ß-Nal, ß-Pal, Trp, Phe, 2,4-dichloro-Phe, pentafluoro-Phe, p-X-Phe, or o-X-Phe, wherein X is CH₃, Cl, Br, F, OH, OCH₃ or NO₂;

A² is Ala, Leu, Ile, Val, Nle, Phe, ß-Nal, pyridyl-Ala, Trp, 2,4-dichloro-Phe, pentafluoro-Phe, o-X-Phe, or p-X-Phe, wherein X is CH₃, Cl, Br, F, OH, OCH₃ or NO₂;

 A^3 is pyridyl-Ala, Trp, Phe, ß-Nal, 2,4-dichloro-Phe, pentafluoro-Phe, o-X-Phe, or p-X-Phe, wherein X is CH_3 , Cl, Br, F, OH, OCH_3 or NO_2 ;

A⁶ is Val, Ala, Leu, Ile, Nle, Thr, Abu, or Ser;

A⁷ is Ala, Leu, Ile, Val, Nle, Phe, ß-Nal, pyridyl-Ala, Trp, 2,4-dichloro-Phe, pentafluoro-Phe, o-X-Phe, or p-X-Phe, wherein X is CH₃, Cl, Br, F, OH, OCH₃ or NO₂;

A⁸ is a D- or L-isomer of Ala, Leu, Ile, Val, Nle, Thr, Ser, Phe, ß-Nal, pyridyl-Ala, Trp, 2,4-dichloro-Phe, pentafluoro-Phe, p-X-Phe, or o-X-Phe, wherein X is CH₃, Cl, Br, F, OH, OCH₃ or NO₂;

each R_1 and R_2 , independently, is H, lower acyl or lower alkyl; and R_3 is OH or NH_2 ; provided that at least one of A^1 and A^8 and one of A^2 and A^7 must be an aromatic amino acid; and further provided that A^1 , A^2 , A^7 and A^8 cannot all be aromatic amino acids.

25 (currently amended): A <u>The</u> method according to claim 24 wherein the linear somatostatin agonist is H-D-Phe-p-chloro-Phe-Tyr-D-Trp-Lys-Thr-Phe-Thr-NH₂, H-D-Phe-p-NO₂-Phe-Tyr-D-Trp-Lys-Val-Phe-Thr-NH₂, H-D-Nal-p-chloro-Phe-Tyr-D-Trp-Lys-Val-Phe-Thr-NH₂, H-D-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-NH₂, H-D-Phe-Phe-Tyr-D-Trp-Lys-Val-Phe-Thr-NH₂, H-D-Phe-Phe-Tyr-D-Trp-Lys-Val-Phe-Thr-NH₂, H-D-Phe-Phe-Tyr-D-Trp-Lys-Val-Phe-Thr-NH₂, H-D-Phe-Ala-Tyr-D-Trp-Lys-Val-Phe-Thr-NH₂.

26 (currently amended): A $\underline{\text{The}}$ method according to claim \pm $\underline{3}$ wherein the somatostatin agonist is

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$$\label{eq:hoch} \text{HO(CH$_2$)$}_2\text{-N} \\ \hline \text{N-(CH$_2$)$}_2\text{-CO-D-Phe-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-NH$}_2 \\ \\ \end{array}$$

$$\label{eq:hoch} \text{HO(CH}_2)_2\text{-N} \qquad \text{N-(CH}_2)_2\text{-SO}_2\text{-D-Phe-Phe-Phe-D-Trp-Lys-Thr-Phe-Thr-NH}_2$$

or

27 (canceled)

28 (currently amended): A \underline{The} method according to claim 3 wherein said patient is obese.

29 (canceled)

30 (currently amended): A $\underline{\text{The}}$ method according to claim 7 wherein said patient is obese.